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Substitute for form 1449/PTO		Complete if Known Application Number 10/785,497 Filing Date February 24, 2004 First Named Inventor Mark W. Becker et al. Art Unit 1657 Examiner Name P.C. Martin Attorney Docket Number 249.P2	
INFORMATION DISCLOSURE STATEMENT BY APPLICANT <i>(Use as many sheets as necessary)</i>			
Sheet	1	of	2

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FOREIGN PATENT DOCUMENTS						
Examiner Initials*	Cite No. ¹	Foreign Patent Document	Publication Date	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages Or Relevant Figure Appear	†
		Country Code ² , Number ³ , Kind Code ⁴ (if known)	MM-DD-YYYY			
	BA	WO-0208241	01-31-2002	Gilead Sciences Inc et al.		
	BB	WO-9200988	01-23-1992	Bodor Nicholas S		

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Sheet	2	2	

NON PATENT LITERATURE DOCUMENTS			
Examiner Initials	Cite No. ¹	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T ²
	CA	Beach et al. (1998) "Chemotherapeutic agents for human immunodeficiency virus infection: Mechanism of action, pharmacokinetics, metabolism, and adverse reactions." <i>Clinical Therapeutics</i> 20(1):2-25	
	CB	CAMP, N.P. et al. (1995) "Synthesis of Peptide Analogues Containing Phosphonamide Methyl Ester Functionality: HIV-1 Protease Inhibitors Possessing Unique Cell Uptake Properties." <i>Bioorganic & Medicinal Chemistry</i> 3(3):297-312	
	CC	Cihlar et al. (2006) "Suppression of HIV-1 Protease Inhibitor Resistance by Phosphonate-mediated solvent anchoring." <i>Journal of Molecular Biology</i> 363(3):635-647	
	CD	Eddershaw et al. (2000) "ADME/PK as part of a rational approach to drug discovery" <i>Drug Discovery Today</i> 5(9):409-414	
	CE	FRANCHETTI, P., et al. (1998) "Potent and Selective Inhibitors of Human Immunodeficiency Virus Protease Structurally Related to L-94,746," <i>Antiviral Chemistry & Chemotherapy</i> 9(4):303-309	
	CF	GULICK (2003) "New Antiviral Drugs," <i>Clinical Microbiology and Infectious Diseases</i> 9:186-193	
	CG	HOLY (2003) "Phosphonothioalkyl Analogs of Nucleotides," <i>Current Pharmaceutical Design</i> 9:2567-2592	
	CH	Hoggard et al. (2002) "Intracellular pharmacology of nucleoside analogues and protease inhibitors: role of transporter molecules," <i>Current Opinion in Infectious Diseases</i> 15(1):3-8	
	CI	Kiso et al. (1999) "Design of small peptidomimetic HIV-1 Protease Inhibitors and Prodrug Forms," 6(5/6):275-281	
	CJ	Krise et al. (1996) "Prodrugs of Phosphates, Phosphonates, and Phosphinates," <i>Advanced Drug Delivery Reviews</i> 19(2):287-310	
	CK	Kubota et al. (1998) "Novel inhibitory effects of gamma-glutamylcysteine ethyl ester against human immunodeficiency virus type 1 production and propagation," <i>Antimicrobial Agents and Chemotherapy</i> 42(5):1200-1206	
	CL	Lee et al. (2002) "In Vivo and In Vitro Characterization of GS 7340, an isopropylalaninyl phenyl ester prodrug of Tenofovir: selective intracellular activation of GS 7340 leads to preferential distribution in lymphatic tissues. 9th Conference of Retroviruses and Opportunistic Infections, Abstract No. 3847	
	CM	Robbins et al. (1998) "Anti-Human Immunodeficiency Virus Activity and Cellular Metabolism of A Potential Prodrug of the Acyclic Nucleoside Phosphonate 9-R-(2-Phosphonomethoxypropyl) Adenine (PMPA), Bis(Isopropoxyloxymethylcarbonyl) PMPA" <i>Antimicrobial Agents and Chemotherapy</i> 42(3):612-617	
	CN	Zimra et al. (2000) "Uptake of pivaloyloxymethyl butyrate into leukemic esterase-catalyzed hydrolysis," <i>Journal of Cancer Research and Clinical Oncology</i> 126(12):693-698	

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¹Applicant's unique citation designation number (optional). ²Applicant is to place a check mark here if English language Translation is attached.